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- (71) Applicant (*for all designated States except US*): **OOO PKF "ATLANTIS-PAK" [RU/RU]; ul. Onuchkina, 72, kh. im. Lenina, Aksaisky raion, Rostovskaya obl., 346700 (RU).**
- (72) Inventors; and
- (75) Inventors/Applicants (*for US only*): **BORODAEV, Sergei Vasilievich [RU/RU]; ul. 2-ya Koltsevaya, 94, Rostov-na-Donu, 344004 (RU). DAVIDENKO, Oleg Vladimirovich [RU/RU]; ul. Pushkinskaya, 176-115, Rostov-na-Donu, 344022 (RU). DAVIDENKO, Alexandr Vladimirovich [RU/RU]; ul. Ostrovskogo, 48-5, Rostov-na-Donu, 344082 (RU). RYZENKO, Sergei Petrovich [RU/RU]; pr. 40 let Pobedy, 93-54, Rostov-na-Donu, 344111 (RU).**
- (74) Agents: **EGOROVA, Galina Borisovna Law Firm "Gorodissky and Partners" ltd. et al.; Gorodissky & Partners, ul. Bolshaya Spasskaya, 25-3, Moscow, 129010 (RU).**
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(54) Title: **POLYMER FILM FOR FOOD PRODUCTS**

(57) Abstract: A polymer film is disclosed for effectively smoking and/or drying products packed therein, which are to be smoked and/or air dried, in particular, smoked cheeses, smoked, uncooked smoked, and air-and-sunshine dried sausages, meat and fish products. This single-layer film is produced on the base matrix of an aliphatic polyamide and/or copolyamide and/or terpolyamide and contains 4.0-50.0 wt. % of a hydrophilic compound which is in a highly dispersed state, wherein the hydrophilic compound forms a disperse phase with a linear size of the domain being 0.1-3.0 Sg(m)m in a direction perpendicular to the plane of the film and either a polymer compound or a low-molecular substance - salt, may be used as such.

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POLYMER FILM FOR FOOD PRODUCTS

Field of the Invention

5 The present invention relates to packaging film which has sufficiently high permeability in respect to smoke substances and/or to water vapors in order to provide the possibility for effective smoking and/or drying products packed therein and intended for smoking and/or air drying, in particular, smoked cheeses, smoked, uncooked smoked sausages and air dried sausages, jerked meat and fish products.

10

Background of the Invention

Traditionally, natural casings of animal intestines, and also synthetic casings on the base of collagen and cellulose, having high steam and smoke permeability, are used as casings for smoked sausage products. However, natural intestine casings are an expensive and scarce product, and furthermore are subject to bacterial spoilage. The production of cellulose and collagen synthetic casings is based on low productive and multistage solution technologies related to the use of such toxic materials as carbon disulfide and formaldehyde and presumes a large amount of harmful liquid wastes. This results in a relatively high cost of such casings.

20

Due to the above-mentioned reason, a desirable aim is the development of synthetic analogues of the described casings, which analogues would be produced by highly productive extrusion methods.

Linear aliphatic polyamides or mixtures thereof with other polymers such as an ionomer resin, modified ethylene vinyl acetate copolymers and/or modified polyolefins are disclosed in EP № 0 139 888, published 8 May 1985 as polymer materials suitable for making smokable casings. Wherein, the glass transition temperature of the polymer material decreases depending on the absorption of water. Under conditions of "wet" smoking the products obtain proper fragrance and taste. These casings readily pass the smoke substances, have excellent mechanical properties and low gas permeability. However, due to low vapor permeability they cannot be used instead of natural, collagen or cellulose casings and they purport obtaining therein a somewhat different, more moist product than

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traditionally smoked sausage.

A casing for smoked food products, suitable for both smoking and for storing therein cured products, i.e. having high barrier properties in respect to oxygen and vapor, is disclosed in US patent № 5 084 310, published 25 January 5 1992. The packaging material comprises at least one layer of a mixture of 5-60 wt. % of polyvinylidene chloride (PVDC) and 95-40 wt. % of polyamide. In order to increase the gas barrier properties of the film, PVDC is introduced into a smokable polyamide. A copolymer, mainly consisting of vinylidene chloride, preferably a copolymer of 65-98 wt. % of vinylidene chloride and 2-35 wt. % of a 10 monomer capable of forming a copolymer with vinylidene chloride, for example, vinyl chloride, (met)acrylic acid, acrylonitrile, is used as the PVDC. A polyamide, having a low melting point, not exceeding 210°C, is used as the polyamide to be mixed with the PVDC. Nevertheless, this casing does not pass the smoke substances to a sufficient degree and has high barrier properties in respect to water 15 vapors. A sensor test on the capability for the products to be smoked in this casing shows fragrance and taste of the smoked product to be approximately at the level of a casing of pure polyamide but significantly worse than in the case of a cellulose casing.

In US patent № 4 851 245, published 25 July 1989, smokable films are 20 produced from a mixture of 50-80% polyamide (PA) with a melting point of 120-210°C and 20-50% of a copolymer of ethylene and vinyl alcohol. Polyamide 6.66, polyamide 612 or mixtures thereof are claimed as the polyamides. The packaging film has good smoke permeability, which in this patent is evaluated according to the permeability in respect to vapors of methanol, but it was not the object of this 25 invention to reduce the moisture permeability of this film, made in the form of a tubular casing. Moreover, in one of the embodiments of the invention the casing comprises a moisture impermeable layer.

A film for smoking and storage is disclosed in EP № 0 252 597, published 13 January 1988. This film is made of a blend of polymers in an amount of 80-98 30 wt. % and additives in an amount of 2-20 wt. %, wherein the polymer mixture consists in turn of 30-100 wt. % of polyamide and 0-70 wt. % of polyolefins. This additive that actually is a plasticizer, is compatible with the polymer components,

dissolves in water and/or oil and is in a liquid state at 70-95°C, which makes it possible for it in the process of successive heating to sweat in the inner and/or outer surface of the film, after which the film becomes vapor and gas impermeable, as if it is made from the same polymer materials, but with no additive. The following are used as an additive: aliphatic alcohols, polyglycols, esters of polyatomic alcohols, esters of dibasic aliphatic acids, esters of polyvalent oxycarboxylic acids, ethers of aliphatic acids and epoxy plasticizers. Impairment of the mechanical properties, which may result in rupture of the film when sausage is being stuffed into the casing, relates to drawbacks of these strongly plasticized films. Furthermore, premature sweating of the plasticizer on the outer surface may result in impairment of the printing ink adhesion, and inside - may promote the acquirement of a foreign taste by the product packed in the film.

The most pertinent prior art is EP № 0 920 808, published 9 June 1999 which discloses a film used for packing cooked processed meat products, such as sausage and ham, and also for cheeses, in the case where they are smoked and subjected to heat treatment. The film is prepared either from acetate-propionate cellulose or a mixture thereof with polyamide with a weight ratio of the first to the second being 5-90 : 95-5, preferably 5-30 : 95-70 with a thickness of from 5 to 90 µm, wherewith the film in accordance with the invention has a permeability in respect to water vapor equal to 300-363 g/m² per day. However, the films described in the examples, which contain polyamide as the main component, have the permeability, which is not more than 300 g/m² per day. The polyamide may be PA 6.66, PA 11, PA 12, PA 612 or a mixture thereof. Furthermore, the film may contain as an additive 0.1-10.0 wt. % of cellulose powder, which enhances its smokability. Smoking is carried out in the presence of water or water vapor. However, the concrete films described in the examples do not have sufficient water vapor permeability so that the effective loss of moisture is ensured and the sausage product is provided with the necessary consistency.

Summary of the Invention

An object of the instant invention is to develop a reliable, inexpensive synthetic food film that has sufficiently high permeability in respect to smock substances and/or water vapors, in order to provide the possibility for the effective

smoking and/or drying of a product packed therein during preparation according to traditional food processing.

Another object of the instant invention is to develop a synthetic food film that has barrier properties in respect to oxygen, at least at the level of the using
5 polyamide matrix under conditions of low humidity.

The next object of the instant invention is to develop a synthetic food film that has good mechanical properties together with the capability of providing quality smoking and/or air drying.

These objects are achieved in that a polymer film for food products
10 comprises a polyamide matrix and a component providing high permeability in respect to smoke substances and/or water vapors, wherein the aforesaid component is a hydrophilic compound in an amount of 4.5-50.0 wt. %, forming in the polyamide matrix a highly dispersed phase with a linear domain size of 0.1-3.0 μm in a direction perpendicular to a plane of the film, and capable of mixing with
15 at least 10 wt. % of water, wherein:

- it is advisable that aliphatic polyamide, copolyamide or terpolyamide be used as the polyamide matrix;

- polyamide 6 and/or copolyamide 6.66 and/or copolyamide 69 and/or copolyamide 612 and/or terpolyamide 6/66.9 and/or terpolyamide 6/66.12 is used
20 as the polyamide matrix;

- polymers selected from homopolymers and/or copolymers of vinylpyrrolidone, vinyl alcohol, alkyloxazoline, polyacrylamide, cellulose ethers, alkylene glycols, polyalkylene oxides, acrylic acid, metacrylic acid, vinyl alcohol ethers, vinyl alcohol esters are used as the hydrophilic compound; the aforesaid
25 homopolymers and/or copolymers may be water-soluble;

- water-soluble low-molecular substances are used as the hydrophilic compound;

- substances selected from inorganic salts, salts with an organic anion and an inorganic cation are used as the water-soluble low molecular substances;

- the film may additionally contain plasticizers and/or dyes and/or pigments and/or antiblocking and/or technological additives;

- the film may be made unoriented;

- the film may be made with single-axis orientation;
- the film may be made with biaxial orientation;
- the polymer film may be made in the form of a seamless tubular casing or packet, wherewith the form given to the polymer film depends on the food products to be placed therein. Thus, in the case of packing fish or solid cheese, it is advisable to use packets, while in the case of producing different forms of smoked sausage or melted cheese - a tubular casing.

Brief Description of the Drawings which Illustrate the Essence of the Invention

Fig.1 shows an external view of the apparatus for determining phenol permeability of a flat sample of film.

Figs. 2-5 show scanned electronic microphotographs of microcuts of film (thickness of cut 5000 Å), held for 2 sec in water and made with the aid of a "DEH 345" penetrating electronic microscope:

Fig.2 shows a cut of a film prepared in accordance with example № 1;

Fig.3 shows a cut of a film prepared in accordance with example № 2;

Fig.4 shows a cut of a film prepared in accordance with example № 3;

Fig.6 shows a cut of a film prepared in accordance with example № 6.

Figs. 5, 7 show an optical microphotograph of a cut (5000 Å) of a prepared film, the microphotograph made with the aid of a "Geneval" Karl Zeiss Jena optical microscope:

Fig.5 shows a cut of a film prepared in accordance with example № 4;

Fig.7 shows a cut of a film prepared in accordance with example № 10.

The data presented in Figs. 2-7 illustrate the morphology of polymer mixtures of prepared films, but in respect to some, as proof of the high phase dispersity, data are provided on the optical properties of the film (opalescence). In example 1, in spite of the direct confirmation of the existence of a separate highly dispersed phase of polyvinyl pyrrolidone (PVP) by electronic microphotographs, opalescence is not observed, and this, obviously, is related to the close values of the indexes of refraction of PVP and PA 6.66.

Detailed Description of the Preferred Embodiments

In the process of working on the instant invention, we unexpectedly discovered a certain relationship between the distribution of the hydrophilic

additive in the polymer matrix and the capability of this mixture to pass water vapors and smoke components. This relationship will be illustrated in the examples below. It was found that if such a compound is physically compatible with the main polymer, just like as if it will form therein a rough dispersion (the
5 size of additive domains is comparable with the thickness of the film and is 10-40 μm), then a significant increase of the vapor and smoke permeability is not observed. Compatibility is understood to mean the capability of two or more components to form liquid or solid systems that are homogeneous on a molecular level. This concept includes both the swellability and solubility. A domain is
10 understood to mean a continuous region of a homogeneous (at the micro level) composition. To the contrary, in the case where the hydrophilic compound forms a highly dispersed phase, a sharp increase of these parameters is observed. Obviously, this effect is related to penetration of the hydrophilic phase domains of a certain size into the compact surface layer of the polymer film, the fact of
15 existence of which is well known to specialists. It is clearly manifested on the presented electronic microphotographs in the form of a dark strip having a thickness of up to 3 μm . The appearance of this layer is usually related to the speed gradient of the flow of the polymer during die and orientation drawing, as a result of which the surface layers of the film are oriented to a greater degree and
20 they have greater density and crystallinity. In the case of polyamide, this layer is "barrier" since it has substantially reduced hydrophily and, consequently, moisture permeability, this determining the moisture permeability of the polyamide film as a whole. Apparently, it is just because of this that in the cases where the hydrophilic compound is compatible with polyamide and where the size of the
25 phase domains of the additive substantially exceeds the thickness of the "barrier" layer and they are wholly positioned outside the "barrier" layer, the "flow" of the polymer material during the stretching bends round a large size defect and in that case a significant changing the vapor and smoke permeability of the film is not observed. And only when the domain of the hydrophilic phase have a size, which
30 is comparable with the thickness of the "barrier" layer, the total permeability does significantly increase. A condition for such dispersity is the close affinity of the hydrophilic compound to polyamide, this affinity being expressed in its optimum

solubility in the polyamide melt. However, the affinity should not be so close that it results in their miscibility (the absence of the separated phase of a hydrophilic compound) in a solid state.

5 In order to enhance the performance of the film, its composition may include plasticizers, dyes, pigments, antiblocking and technological additives. Packaging films may be oriented and unoriented. They may be bent by stretching or setting, if it is necessary to obtain sausage with a certain curvature.

10 Forming the polymer mixture into a tubular casing may be carried out by any of the known methods, for example, extrusion through an annular die directly into a tubular casing or by extrusion through a flat-slit die and obtaining first a flat sheet and subsequently preparing a tubular casing therefrom.

During the production of a packet, a section of a tubular film of the necessary length is sealed at one end.

15 Films produced by the facile one-stage processing of polymer feedstock have excellent mechanical properties, which make it possible to give a shape to the products packed therein and to maintain that shape in the process of production and storage. Furthermore, no rupture of a film produced in the form of a seamless tubular casing occurs when it is stuffed under pressure.

20 The proposed film is suitable for packing and subsequently smoking cheese, smoking and/or air drying meat and fish products during their production in accordance with traditional technologies. It makes it possible to prepare products with the required remaining content of moisture for the time provided by the technology of production of the said food products. Air drying is understood to mean a loss of moisture in the process of preparing the product.

25 Due to the fact that the proposed film in a moist state becomes gas permeable, it may be used during the preparation of uncooked smoked sausages, the process of preparation of which includes fermentation which requires aerobic conditions and is accompanied by the release of CO₂. After the curing process is over, when the humidity of the sausage-meat is reduced to low values, the film
30 will again acquire gas-barrier properties.

The packaging film produced in the form of a packet may be used for products, the preparation of which presumes their being cut and subsequently

dried at room or at a higher temperature, for example, dried fruits.

In respect to the properties of the film of the present invention, it does not make any difference in which form—flat sheet, tubular casing or packet—it is produced.

5 The smoke permeability of a film was evaluated as the permeability in respect to phenol - one of the main constituents of the smoke components. The extraction method of determining the phenol permeability of sausage casings is described below.

10 Phenol permeability is characterized by the concentration of phenol, which has passed through the casing from a standard solution into an aqueous extract under predetermined conditions.

15 Determination of the phenol permeability of a flat sample of film is carried out with the aid of a special apparatus (Fig. 1), which is a unit consisting of two chambers divided by the casing, a KFK-2 photo-electrocalorimeter and a pH-meter.

20 The apparatus is placed in a drying cabinet preliminarily heated to 85°C, where it is held for 80 min, wherein there is a solution for determining phenol permeability in the lower chamber, 100 ml of distilled water in the upper. The concentration of the phenol is determined in the solution from the upper chamber of the unit, wherefore it is extracted with butyl acetate. The standard solution of phenol is prepared by the dissolution of 0.1 g of freshly-distilled crystalline phenol in 50 ml of ethyl alcohol. The solution for determining the phenol permeability is prepared from 10 ml of a standard solution of phenol with the addition of 1 ml of acetic acid and with the volume brought to 100 ml with
25 distilled water. The weight concentration of phenol in the sample being analyzed is found by a preliminarily plotted calibration curve. The calibration relationship is plotted in the coordinates: weight concentration of phenol, $\mu\text{g/ml}$ - optical density of the sample minus the optical density of a blank experiment. The optical density of the extract is measured in vessels 5 cm long with a
30 photoelectrocolorimeter at $\lambda = 490 \text{ nm}$ in respect to a pure solvent.

Preferable embodiments of the invention are shown in the following examples.

Example 1

A mixture comprising 94 wt. % of granulated PA 6.66 ("Ultramid C35" BASF brand) and 10 wt. % of polyvinyl pyrrolidone (trademark "Povidon" commercially available from BASF) is loaded into an extruder and melted. The melt with a temperature of 230-235°C is extruded through an annular die and a primary tube is formed. Then at a temperatures of 60°C it is subjected to biaxially-orientational drawing with air-blowing (double-bubble process), wherein the coefficient of longitudinal stretch is 2.6, lateral - 3.0. After that, the tubular film having a caliber of 24 mm and a thickness of 19-21 µm is subjected to relaxation annealing at a temperature of 160-180°C for 15 sec, cooled to 20°C and wound into a roll.

Spaces with a cross section of 0.1-0.2 µm, formed at the place of the water-soluble phase, are clearly seen in Fig. 2.

Then the tubular casing is used in the production of two kinds of cured sausage.

Method 1. Preparation of semi-smoked sausages

The casing described above is packed with the sausage stuffing for the semi-smoked sausages. The prepared sausage is held (set) for 24 hours at a temperature of 3°C, dried and browned for 1 hour at a temperature of 95°C. Then it is smoked with the smoke of smoldering birch sawdust for 5 hours at a temperature of 45°C and relative humidity of 95%, after which it is dried at a temperature of 11°C for 36 hours.

Method 2. Preparation of uncooked smoked sausages

The casing is packed with sausage stuffing for uncooked smoked sausages. The prepared sausage is held in a ripening chamber for six days at a temperature of from 24°C to 18°C and a relative humidity which gradually decreases during that period from 98 to 90% in order to carry out fermentation of the stuffing. Then it is smoked with the smoke of smoldering birch sawdust for 5 hours at a temperature of 20-25°C and relative humidity of 85%, after which it is dried at a temperature of 15-12°C, gradually reducing the relative humidity from 85 to 74% during 25 days.

Polymer films, prepared in accordance with the examples presented below

were used for the preparation of semi-smoked sausages according to method 1 and uncooked smoked sausages according to method 2.

Example 2

The casing is produced in accordance with example 1, wherewith 4 wt. %
5 of polyvinyl pyrrolidone is introduced as the hydrophilic compound.

A continuous medium, which does not contain any voids, is clearly seen in Fig. 3.

Example 3

The casing is produced in accordance with example 1, wherewith
10 granulated PA 6 ("Ultramid B35") in an amount of 80 wt. % is used as the polyamide base matrix, the hydrophilic compound is 15 wt. % of polyvinyl alcohol (trademark "Mowiol 5-88" commercially available from Clariant) with a degree of saponification 88% and viscosity of a 4% aqueous solution being 5 centipoises, and a plasticizer - glycerin in an amount of 5 wt. %.

15 The obtained casing has an expressed opalescence, intensifying when moistened, which indicates the presence therein of a separate phase that is in a highly dispersed state. Voids having a cross-sectional size of 0.2-3.0 μm are clearly seen in Fig.4, these spaces being formed at the point of water-soluble phase.

Example 4

20 The casing is produced in accordance with example 3, wherein polyvinyl alcohol (trademark "Mowiol 6-98" commercially available from Clariant) is used with a hydrolysis degree of 98% and with viscosity of a 4% aqueous solution being 6 centipoises.

25 The casing has a lusterless appearance with marked rough inclusions. Roughly dispersed PVA domains with a size of 4-10 μm are clearly seen in Fig.5. A general thickening of the shell is evident near especially large domains.

Example 5

The casing is produced in accordance with example 1, wherein a polymer
30 mixture is used which consists of 75 wt. % of granulated PA 6.66 ("Ultramid C35" BASF brand) and 25 wt. % of poly N-ethyl oxazoline with a molecular weight of 500000 (trademark "Aquazol-500" commercially available from

Polymer Chemistry Innovations Inc.).

The obtained casing has an expressed opalescence, intensifying when moistened, which indicates the presence therein of a physical heterogeneity of the polymer mixture and, accordingly, its highly dispersed state.

5

Example 6

The casing is produced in accordance with example 5, wherein 15 wt. % of poly N-ethyl oxazoline, "Aquazol-500", is used as the hydrophilic compound.

A polymer medium which does not have any empty spaces is shown in Fig.6.

10

The prepared casing appears optically homogeneous, opalescence is not manifested even with moistening.

Example 7

The casing is produced in accordance with example 1, wherein the polymer mixture consists of a mixture of 90 wt. % of granulated PA 6.66 ("Ultramid C35") and 10 wt. % of a mixture of sodium acetate/potassium acetate powders (40:60, $T_{\text{melt}} \sim 180^{\circ}\text{C}$).

15

The obtained casing has an expressed opalescence, intensifying with moistening, which indicates the presence therein of a separate phase which is in a highly dispersed state.

20

Example 8

The casing is produced in accordance with example 1, using wherein a polymer mixture consisting of 90 wt. % of granulated PA 6.66 ("Ultramid C35") and 10 wt. % of a mixture of sodium metaphosphate/potassium metaphosphate powders (50:50, $T_{\text{melt}} \sim 150^{\circ}\text{C}$).

25

The obtained casing has an expressed opalescence, intensifying with moistening, which indicates the presence therein of a separate phase which is in a highly dispersed state.

Example 9

The casing is produced in accordance with example 1, wherein the polymer mixture consists of 90 wt. % of granulated PA 6.66 ("Ultramid C35") and 10 wt. % of calcium chloride, compatible with a polyamide melt (see, for example, "Polyamides" edited by M. Kohan, N.Y., 1995, p. 439).

30

The casing has marked opalescence.

Example 10

The casing is produced in accordance with example 1, wherein the polymer mixture is prepared from 90 wt. % of granulated PA 6.66 ("Ultramid C35") and 10 wt. % of sodium chloride powder, which is not compatible with polyamide in a viscous-flow and solid state.

The prepared casing has an expressed dullness and contains rough inclusions. Roughly dispersed NaCl crystals having a size up to 15 μm are clearly seen in Fig.7. A general thickening of the casing is evident near especially large domains.

Example 11

The casing is produced in accordance with the technology of example 1 from polyamide 6.66 ("Ultramid C35") without the addition of a hydrophilic compound.

The characteristics of all the produced films and the data in respect to their capability for smoking are summed up in the Table, wherein:

¹ Mechanical characteristics of the tubular casings were determined on a "Shimadzu AGS-H" autograph with a speed of moving the gripping devices apart equal to 100 mm/min.

² Water vapor permeation rate is determined at 30°C and relative humidity of 65% in accordance with DIN 53 122-74.

³ Phenol vapor permeation rate is determined at 85°C on an apparatus described in accordance with the Method.

⁴ Oxygen permeation rate is determined at 30°C and relative humidity of 65 % in accordance with DIN 53 380-69.

⁵ ++ Presence of a tough dark crust of coagulated protein with a thickness of 0.5-1 mm, easily separated from the casing on the surface of sausage.

+ Presence of a dark crust with a thickness less than 0.5 mm, which can be separated from the casing.

- No dark crust present, loaf which is not swollen.

-- Swelling of casing during the stage of fermentation.

⁶ ++ Clear fragrance and taste of smoking.

- + Weak fragrance of smoking.
- No fragrance of smoking.
- ⁷ O - homogeneous.
- RD - roughly dispersed.
- 5 HD - highly dispersed.

10 The data presented in the table clearly confirm the fact that films prepared in accordance with examples 1, 3, 5, 7-9, and containing a hydrophilic phase with the size of the domains within the range of 0.1-3.0 μm have higher permeability in respect to the smoke substances and/or water vapors as compared with films prepared in accordance with examples 2, 4, 6, 10, and ensure during smoking a good appearance and good taste of the sausages, while maintaining high mechanical properties.

Characteristics	Examples											"Cutesin" thickness 60 μm
	1	2	3	4	5	6	7	8	9	10	11	
1	2	3	4	5	6	7	8	9	10	11	12	13
Tensile strength, MPa ¹	151	155	164	162	148	153	169	170	163	171	167	-
Elongation to break, %	155	131	123	119	190	186	133	137	137	100	110	-
Phenol vapor permeation rate, g/m ² day ³	150	97	131	103	162	107	120	118	130	101	99	160
Water vapor permeation rate, g/m ² day ²	453	215	509	250	480	225	463	495	515	211	216	1200
Oxygen permeation rate, cm ³ /m ² atm. Day ⁴ at 25°C: at relative humidity 60% at relative humidity 95%	29 120	8 48	17 115	21 60	21 126	30 63	28 110	25 279	18 284	20 55	27 54	800 950

1	2	3	4	5	6	7	8	9	10	11	12	13
Loss of mass during smoking processing of sausage, %: With method 1 With method 2	10.6	6.2	12.8	5.1	9.8	6.0	10.2	9.9	10.1	5.8	5.3	13.5
	11.2	8.1	14.5	6.3	12.5	7.5	13.0	12.1	14.0	6.8	6.3	15.5
Appearance of sausage Stick ⁵ , made according to: Method 1 Method 2	++	-	++	+	++	-	+	+	++	-	-	++
	++	--	++	+	++	--	+	+	++	--	--	++
Smoking fragrance test ⁶ Made according to Method 1 Method 2	++	-	++	+	++	-	++	++	++	-	-	++
	++	-	++	+	++	-	++	++	++	-	-	++
Morphology of hydrophilic additive	HD	O	HD	RD	HD	O	HD	HD	HD	RD	-	-

CLAIMS

1. A polymer film for food products, comprising a polyamide matrix and a component providing high permeability in respect to smoke substances and/or water vapors, wherein said component is a hydrophilic compound in an amount of 4.5-50.0 wt. %, forming in the polyamide base matrix a highly dispersed phase with a linear domain size of 0.1-3.0 μm in a direction perpendicular to a plane of the film, and capable of mixing with at least 10 wt. % of water.

2. A polymer film for food products according to claim 1, wherein said aliphatic polyamide and/or copolyamide and/or terpolyamide is used as the polyamide matrix.

3. A polymer film for food products according to claim 2, wherein said polyamide 6 and/or copolyamide 6.66 and/or copolyamide 69 and/or copolyamide 612 and/or terpolyamide 6/66.9 and/or terpolyamide 6/66.12 is used as the polyamide matrix;

4. A polymer film for food products according to any one of claims 1-3, wherein said polymers selected from homopolymers and/or copolymers of vinylpyrrolidone, vinyl alcohol, alkyloxazoline, alkylene glycols, acrylamide, alkylene oxides, acrylic acid, metacrylic acid, maleic anhydride, vinyl alcohol ethers, vinyl alcohol esters, and also cellulose ethers are used as the hydrophilic compound.

5. A polymer film for food products according to claim 4, wherein among said homopolymers and/or copolymers, those are used which are water-soluble;

6. A polymer film for food products according to any one of claims 1-3, wherein said water-soluble low-molecular substances are used as the hydrophilic compound.

7. A polymer film for food products according to claim 6, wherein said substances selected from inorganic salts, salts with an organic anion and an inorganic cation are used as the water-soluble low-molecular substances;

8. A polymer film for food products according to any one of claims 1-7, wherein said film includes plasticizers and/or dyes and/or pigments and/or antiblocking and/or technological additives.

9. A polymer film for food products according to any one of claims 1-8, wherein said film is made unoriented;

10. A polymer film for food products according to any one of claims 1-8, wherein said film is made with single-axis orientation.

5 11. A polymer film for food products according to any one of claims 1-8, wherein said film is made with biaxial orientation;

12. A packaging for a polymer film for food products, which packaging is a tubular casing or packet, wherein it is made of a polymer film according to any one of claims 1-11.

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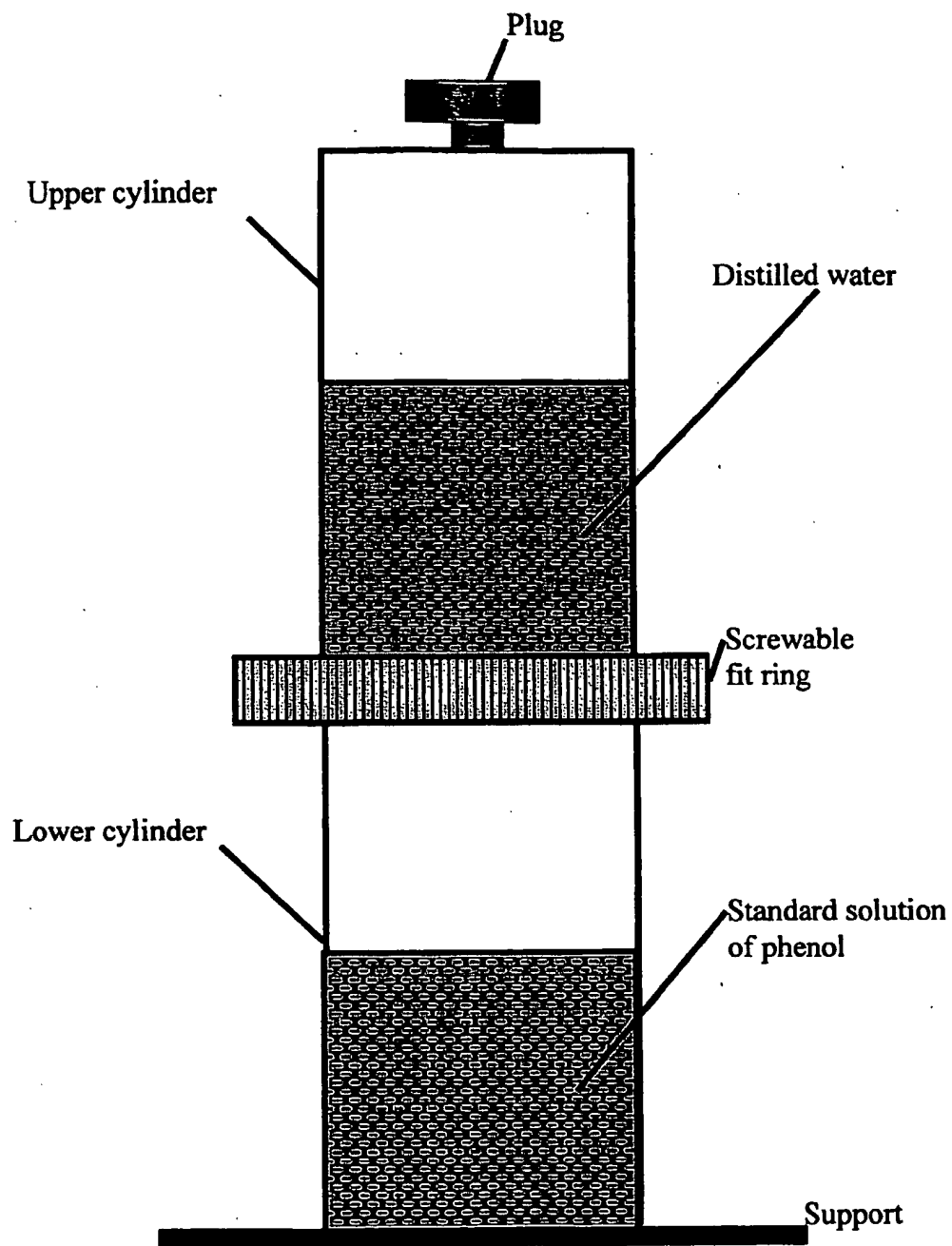


Fig. 1

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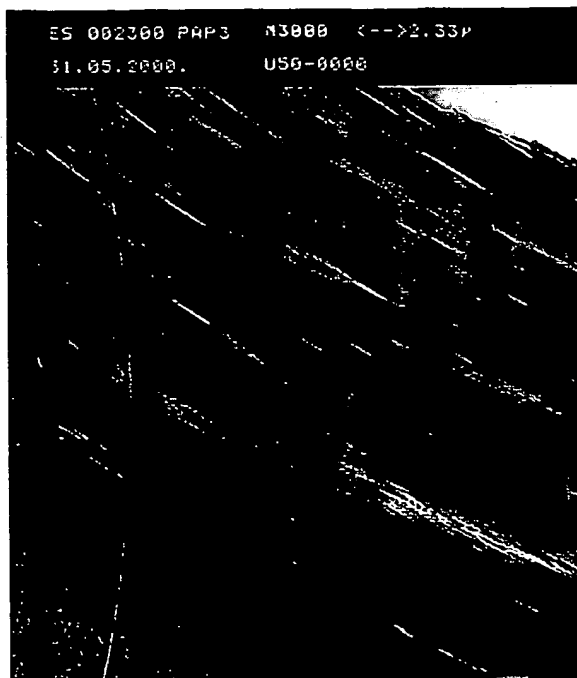


Fig. 2

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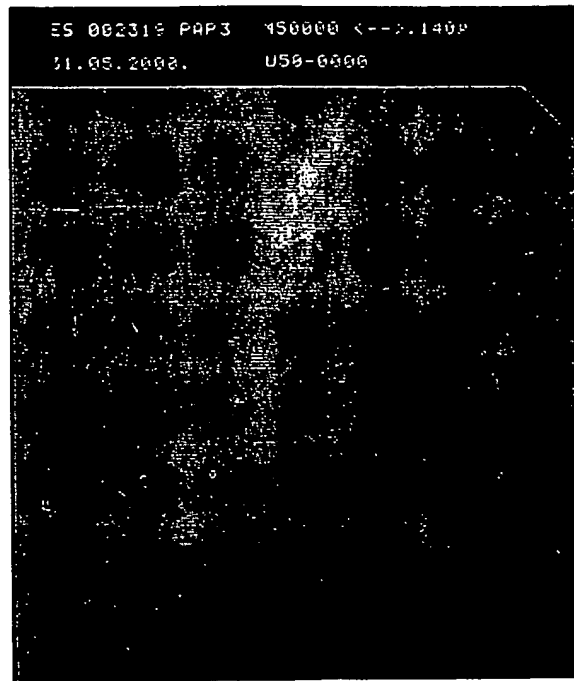


Fig. 3

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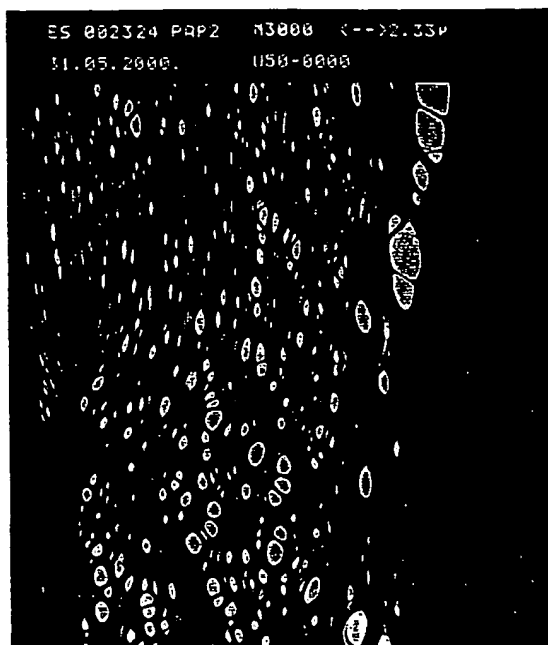


Fig. 4

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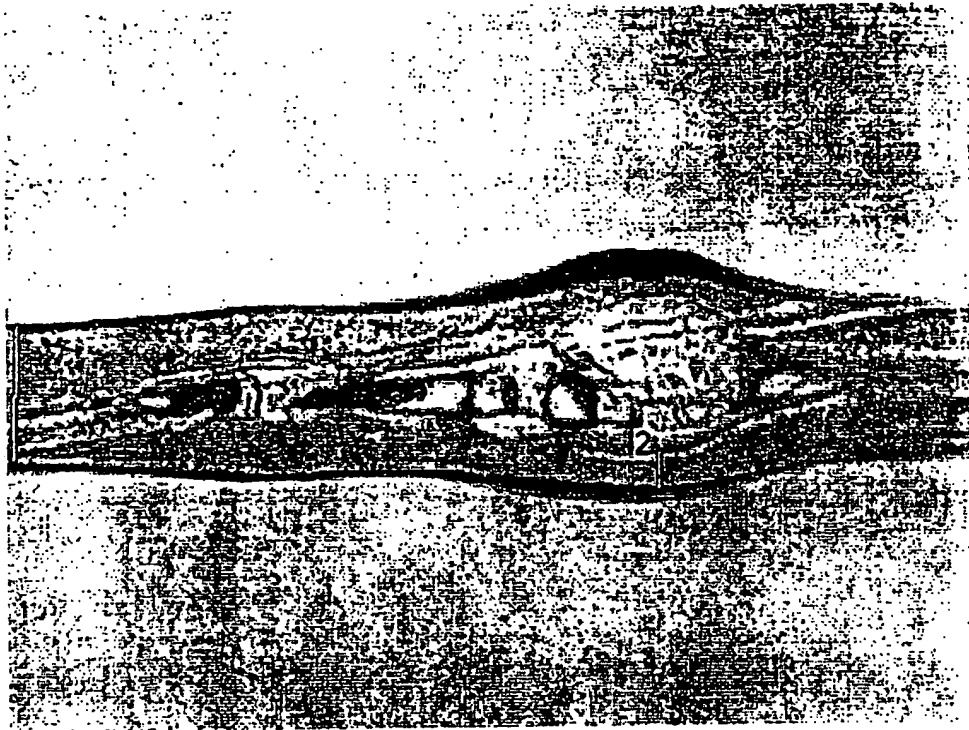


Fig. 5

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*Fig. 6*

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INTERNATIONAL SEARCH REPORT

International Application No.
PCT/RU 02/00115

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A22C13/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 A22C C08J

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

PAJ, WPI Data, EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DE 27 24 252 A (HOECHST AG) 30 November 1978 (1978-11-30) page 10, line 10 - line 20 page 11, line 2 -page 12, line 7; claims 1-18	1-6, 12
Y	EP 0 994 168 A (NITTO DENKO CORPORATION) 19 April 2000 (2000-04-19) page 4, line 5 - line 37 page 4, line 50 -page 5, line 1 page 5, line 25 -page 6, line 8; claims 1-12	1-12
Y	DE 17 17 062 B (SOCIETE CIVILE DES PRODUITS LIFINE) 18 June 1970 (1970-06-18) column 2, line 51 -column 3, line 43; claim 1	1-12

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

20 August 2002

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Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

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Permentier, W

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	EP 0 139 888 A (NATURIN-WERK BECKER & CO.) 8 May 1985 (1985-05-08) cited in the application claims 1-20	1-12
A	EP 0 640 289 A (WOLFF WALSRÖDE AG) 1 March 1995 (1995-03-01) claims 1-9	1
A	EP 0 920 808 A (GUNZE KOBUNSHI CORPORATION) 9 June 1999 (1999-06-09) cited in the application claims 1-10	1
A	EP 0 005 765 A (HOECHST AG) 12 December 1979 (1979-12-12) page 5, line 7 - line 22; claims 1-10	1

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/RU 02/00115

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
DE 2724252	A	30-11-1978	DE 2724252 A1	30-11-1978
			AT 372394 B	26-09-1983
			AT 386078 A	15-02-1983
			CH 635294 A5	31-03-1983
			FR 2401073 A1	23-03-1979
			GB 1603014 A	18-11-1981
			US 4243074 A	06-01-1981
EP 994168	A	19-04-2000	JP 2000119410 A	25-04-2000
			EP 0994168 A2	19-04-2000
			US 6372339 B1	16-04-2002
			JP 2000281806 A	10-10-2000
DE 1717062	B	18-06-1970	CH 403455 A	30-11-1965
			DE 1717062 B1	18-06-1970
			FR 1260250 A	05-05-1961
			GB 917965 A	13-02-1963
			LU 37997 A	
			NL 122082 C	
			NL 258156 A	
			US 3329509 A	04-07-1967
EP 139888	A	08-05-1985	DE 3426723 A1	21-02-1985
			AT 31004 T	15-12-1987
			AU 567179 B2	12-11-1987
			AU 3126784 A	14-02-1985
			CA 1235018 A1	12-04-1988
			DE 3467684 D1	07-01-1988
			DK 365084 A	29-01-1985
			EP 0139888 A1	08-05-1985
			ES 287175 U	16-12-1985
			ES 534687 D0	16-10-1985
			ES 8600873 A1	16-02-1986
			FI 842965 A	29-01-1985
			JP 1952532 C	28-07-1995
			JP 6073428 B	21-09-1994
			JP 60105448 A	10-06-1985
			ZA 8405811 A	26-03-1986
EP 640289	A	01-03-1995	DE 4328517 A1	02-03-1995
			AT 179573 T	15-05-1999
			DE 59408204 D1	10-06-1999
			DK 640289 T3	01-11-1999
			EP 0640289 A2	01-03-1995
			ES 2131133 T3	16-07-1999
			JP 7075482 A	20-03-1995
EP 920808	A	09-06-1999	JP 11164648 A	22-06-1999
			AU 9518198 A	24-06-1999
			EP 0920808 A1	09-06-1999
			NO 985596 A	07-06-1999
			ZA 9811111 A	08-06-1999
EP 5765	A	12-12-1979	DE 2822886 A1	29-11-1979
			BR 7903267 A	11-12-1979
			CA 1129729 A1	17-08-1982
			DE 2961355 D1	21-01-1982
			EP 0005765 A1	12-12-1979

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/RU 02/00115

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 5765	A	FI 791648 A , B, JP 1509377 C JP 54156786 A JP 63058540 B US 4289171 A	27-11-1979 26-07-1989 11-12-1979 16-11-1988 15-09-1981